interns for the continuing NIDA Summer Research Internship Program. This request is to allow NIDA to collect information from applicants in order to meet the goals of the program and IC mission. Applicant eligibility for this program was 17 years, but is now open to those 18 and over in the year of application per NIH policy document 2019 High School Summer Internship Program (HS–SIP) Policy. NIDA will request clearance for any additional forms should new programs be introduced in the future. The information ensures that students applying to this program meet basic eligibility requirements; indicates their interest in substance abuse research, future career goals, and, if selected for the program, what research they prefer to conduct. The information also enables decision-making regarding which applicants will be selected for internships. In each case, completing the application is voluntary, but in order to receive due consideration, the prospective applicant must complete all fields required by the program.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 250.

ESTIMATED ANNUALIZED BURDEN HOURS

Form	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Estimated total annual burden hours
Summer Internship	Individuals-household	250	1	1	250
Total			250		250

Dated: October 16, 2019.

Lanette A. Palmquist,

Project Clearance Liaison, National Institute on Drug Abuse, National Institutes of Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Advancing Translational Sciences; Notice of Workshop

Notice is hereby given of a workshop convened by the National Center for Advancing Translational Sciences Cures Acceleration Network Review Board.

Purpose

The National Center for Advancing Translational Sciences is hosting the Cures Acceleration Network Review Board (CAN RB) Workshop: The CAN RB advises and provides recommendations to the NCATS Director with respect to significant barriers to successful translation of basic science into clinical application. In support of this mandate, the CAN RB will co-host this public workshop to discuss challenges around finding new uses for drugs that are already on the market but lack commercial and regulatory incentives for research and development.

The Workshop is being co-sponsored by the NCATS Cures Acceleration Network Review Board, NCATS Drug Development Partnership Programs, Food and Drug Administration, and Reagan-Udall Foundation for the FDA.

Name of Committee: National Center for Advancing Translational Sciences Cures Acceleration Network Review Board.

Type of Meeting: Repurposing Off-Patent Drugs: Research & Regulatory Challenges.

Date: December 5–6, 2019.

Time: 7:30 a.m. to 5:00 p.m., Eastern Standard Time (EST).

Agenda: The Workshop will assess challenges around finding new uses for drugs that are already on the market but lack commercial and regulatory incentives for research and development. On December 5, we will map out the challenges to repurposing off-patent drugs. On December 6, we will host interactive work sessions focused on capturing possible solutions.

Place: Hilton Washington DC/Rockville Hotel & Executive Meeting Center, 1750 Rockville Pike, Rockville, MD 20852.

Cost: The meeting is free and open to the public.

Registration: Registration is required. Using the link below, attendees can register by December 2, 2019, 5:00 p.m. EST. Early registration is recommended due to limited seating. *https://reaganudall.salsalabs.org/repurposingoffpatentdrugsworkshop/index.html.*

Access: Twinbrook Metro Station (Red Line).

Contact Person: Bobbie Ann Mount, Ph.D., Program Officer, Drug Development Partnership Programs, Office of the Director, National Center for Advancing Translational Sciences, National Institutes of Health. *Telephone:* 301.435.0824, Email: *NewTherapeuticUses@mail.nih.gov.*

Disability Accommodations: Individuals whose full participation in the workshop will require special accommodations (*e.g.,* sign language, or interpreting services, etc.) must submit a request to the Contact Person listed on the notice at least ten (10) business days prior to the meeting. Such requests should include a detailed description of the accommodation needed and a way to contact the requester if more information is needed to fill the request. Special requests should be made as early as possible. Last minute requests may be made but may not be possible to accommodate.

Security: Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit. Also, as a part of security procedures, attendees should be prepared to present a photo ID at the meeting registration desk during the check-in process. Pre-registration is recommended. Seating will be limited to the room capacity and seats will be on a first come, first serve basis, with expedited check-in for those who are pre-registered.

Meeting schedule subject to change.

For more information, visit: *https://reaganudall.salsalabs.org/repurposingoffpatentdrugsworkshop/index.html.*

Dated: October 21, 2019.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

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